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**PATENT** 

Attorney Docket No.: A-2326-2/RFT/JJD

### IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

OCT 3 0 2000

BADEMA

In re application of:

KEYT et al.

Serial No. 09/346,069

Filed: July 1, 1999

**NUCLEIC ACIDS ENCODING** For:

> VARIANTS OF VASCULAR **ENDOTHELIAL CELL GROWTH**

**FACTOR** 

Examiner: Unknown

RECEIVED

Group Art Unit: 1646

NOV 0 1 2000

TECH CENTER 1600/2900

CERTIFICATE OF MAILING
Thereby certify that this correspondence, including listed enclosures, is being deposited with the United States Postal Service as First Class Mail in an envelope addressed to: Assistant Commissioner for Patents, Washington, DC 20231 on:

### TRANSMITTAL OF NOTICE TO COMPLY

**Assistant Commissioner** for Patents Washington, DC 20231

Sir:

In response to the Notice to Comply with Requirements for Patent Applications Containing Nucleotide Sequence and/or Amino Acid Sequence Disclosures dated September 29, 2000, enclosed are the following documents:

- 1. Copy of Communication and Notice to Comply;
- 2. Paper copy of SEQUENCE LISTING;

**Serial No.**: 09/346,069 **Filed**: July 1, 1999

- 3. Amendment Re: Sequence Listing
- 4. Computer readable copy of Sequence Listing.

While no fee is currently believed to be due, the Commissioner is authorized to charge any additional fees, including any extension fees, which may be required, or credit any overpayment to Deposit Account No. 06-1300 (Our Order No. A-62326-2/RFT/JJD).

Respectfully submitted,

FLEHR HOHBACH TEST ALBRITTON & HERBERT LLP

Dated: /C

Richard F. Trecartin

Reg. 31,801

Four Embarcadero Center

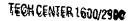
**Suite 3400** 

San Francisco, CA 94111-4187

Telephone: (415) 781-1989

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### DEPARTMENT OF COMMERCE

#### **Patent and Trademark Office**

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Washington, D.C. 20231

ATTORNEY DOCKET NO. FIRST NAMED INVENTOR APPLICATION NO. **FILING DATE** 97/01/99 E A-762326-27R 09/346,069 KEYT **EXAMINER** HM12/0929 KAUFMAN, C FLEHR HOHBACH TEST ALBRITTON & HERBERT LLP PAPER NUMBER **ART UNIT** SUITE 3400 FOUR EMBARCADERO CENTER .646 SAN FRANCISCO CA 94111-4167 **DATE MAILED:** 09729700 OCT 3 0 200

Please find below and/or attached an Office communication concerning this application or proceeding.

**Commissioner of Patents and Trademarks** 

File A-62326-2 Atty DAL 



COMMISSIONER FOR PATENTS
UNITED STATES PATENT AND TRADEMARK OFFICE
WASHINGTON, DC 20231
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| APPLICATION NO./<br>CONTROL NO. | FILING DATE | FIRST NAMED INVENTOR / PATENT IN REEXAMINATION |            | ATTORNEY DOCKET NO. |    |
|---------------------------------|-------------|--|------------|---------------------|----|
|                                 |             |  | EXAMINER   |                     |    |
|                                 |             | IPE  | ART UNIT   | PAPER               |    |
|                                 |             | OCT 3 0 2000 JU                                | DATE MAILE | <b>725</b>          | Cu |

Please find below and/or attached an Office communication concerning this application or proceeding.

#### **Commissioner of Patents**

This application contains sequence disclosures that are encompassed by the definitions for nucleotide and/or amino acid sequences set forth in 37 C.F.R. § 1.821(a)(1) and (a)(2). However, this application fails to comply with the requirements of 37 C.F.R. §§ 1.821-1.825 for the reason(s) set forth on the attached Notice To Comply With Requirements For Patent Applications Containing Nucleotide Sequence And/Or Amino Acid Sequence Disclosures. Applicant must comply with the requirements of the sequence rules (37 CFR 1.821 - 1.825) before the application can be examined under 35 U.S.C. §§ 131 and 132.

No CRF or paper copy of the Sequence Listing was provided as required. When a sequence is presented in a drawing, regardless of the format or the manner of presentation of that sequence in the drawing, the sequence must still be included in the Sequence Listing and a sequence identifier ("SEQ ID NO:X") must be used either in the drawing or in the Brief Description of the Drawings. See MPEP § 2422.02. Sequence identifiers must also be used when a sequence is shown or referred to in the specification or claims. In the instant application, a sequence identifier must be used for the sequences appearing in Figures (e.g., Fig. 1A), as well as in the specification (e.g., p. 44 and 56).

APPLICANT IS GIVEN 30 days FROM THE DATE OF THIS LETTER WITHIN WHICH TO COMPLY WITH THE SEQUENCE RULES, 37 C.F.R.. §§ 1.821-1.825. Failure to comply with these requirements will result in ABANDONMENT of the application under 37 C.F.R. § 1.821(g). Extensions of time may be obtained by filing a petition accompanied by the extension fee under the provisions of 37 C.F.R. § 1.136. In no case may an applicant extend the period for response beyond the six month statutory period. Direct the response to the undersigned. Applicant is requested to return a copy of the attached Notice to Comply with the response.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Claire Kaufman whose telephone number is (703) 305-5791. The examiner can normally be reached Monday-Thursday from 8:30-12:30.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Yvonne Eyler can be reached on (703) 308-6564. The fax number for the organization where this application or proceeding is assigned is (703) 308-4242.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (703) 308-0196.

Claire M. Kaufman

Claire M. Kaufman

Patent Examiner, AU 1646

## **Notice to Comply**

Application No. 09/346.069

Applicant(s)

**KEYT ET AL Art Unit** 

Claire M. Kaufman

# NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

Examiner

Applicant must file the items indicated below within the time period set the Office action to which the Notice is attached to avoid abandonment under 35 U.S.C. § 133 (extensions of time may be obtained under the provisions of 37 CFR 1.136(a)).

The nucleotide and/or amino acid sequence disclosure contained in this application does not comply with the requirements for such a disclosure as set forth in 37 C.F.R. 1.821 - 1.825 for the following reason(s):

- ☑ 1. This application clearly fails to comply with the requirements of 37 C.F.R. 1.821-1.825. Applicant's attention is directed to the final rulemaking notice published at 55 FR 18230 (May 1, 1990), and 1114 OG 29 (May 15, 1990). If the effective filing date is on or after July 1, 1998, see the final rulemaking notice published at 63 FR 29620 (June 1, 1998) and 1211 OG 82 (June 23, 1998).
- 2. This application does not contain, as a separate part of the disclosure on paper copy, a "Sequence Listing" as required by 37 C.F.R. 1.821(c).
- 3. A copy of the "Sequence Listing" in computer readable form has not been submitted as required by 37 C.F.R. 1.821(e).
- 4. A copy of the "Sequence Listing" in computer readable form has been submitted. However, the content of the computer readable form does not comply with the requirements of 37 C.F.R. 1.822 and/or 1.823, as indicated on the attached copy of the marked -up "Raw Sequence Listing."
- ☐ 5. The computer readable form that has been filed with this application has been found to be damaged and/or unreadable as indicated on the attached CRF Diskette Problem Report. A Substitute computer readable form must be submitted as required by 37 C.F.R. 1.825(d).
- 6. The paper copy of the "Sequence Listing" is not the same as the computer readable from of the "Sequence Listing" as required by 37 C.F.R. 1.821(e).
- 7. Other:

Applicant Must Provide:

- An initial or substitute computer readable form (CRF) copy of the "Sequence Listing".
- An initial or substitute paper copy of the "Sequence Listing", as well as an amendment directing its entry into the specification.
- A statement that the content of the paper and computer readable copies are the same and, where applicable, include no new matter, as required by 37 C.F.R. 1.821(e) or 1.821(f) or 1.821(g) or 1.825(b) or 1.825(d).

For questions regarding compliance to these requirements, please contact:

For Rules Interpretation, call (703) 308-4216

For CRF Submission Help, call (703) 308-4212

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